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               CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
               GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
               LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
               PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ,
          UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
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              BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
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                                               EP 2003-700335 20030107
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Preparation of cyclic urea derivatives with 5-HT2c receptor activity

L26 ANSWER 31 OF 99 MARPAT COPYRIGHT 2005 ACS on STN DUPLICATE 6

Bromidge, Steven Mark; Lovell, Peter John; Goodacre, Caroline

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AB Title compds. I [p = 0-5; m = 1-3; Y = N, C; A = 0, N, CONH, NHCO, etc.; R1 = halo, alkyl, alkoxy, alkylthio, etc.; R2 = H, halo, alkyl, alkoxy, haloalkyl, haloalkyl, haloalkoxy; R3 = amino; X = CH2, CO] are prepared For instance, 2-(3-fluorophenylamino)ethanol (preparation given) is reacted with Mscl/cH2Cl2 followed by 3-benzyloxy-4-methoxyphenyllamine to give the corresponding substituted diamine. This intermediate is treated with phosgene to give 1-(3-benzyloxy-4-methoxyphenyl)-3-(3-fluorophenyl)imidazolidin-2-one. Substitution of this using 1-(2-chloroethyl)piperidineHCl (MsCCH2CH2CMe, K2CO3, reflux, 5 h) afforded II. I exhibit 5-HTZc receptor activity and are useful for the treatment of CNS disorders such as depression or anxiety.

MSTR 1

G1 = CN / CF3 G3 = 45-1 46-3

H2C-4810

G10 = C(O) MPL: claim 1